Page 2 of 14

Amendment and Response Serial No.: 10/780,150

Confirmation No.: 1273 Filed: February 17, 2004

For: REGULATION OF T CELL-MEDIATED IMMUNITY BY D ISOMERS OF INHIBITORS OF

INDOLEAMINE-2,3-DIOXYGENASE

Amendments to the Claims

This listing of claims replaces all prior versions, and listings, of claims in the aboveidentified application:

- 1. (Withdrawn/currently amended) A method of augmenting rejection of <u>tumor</u> cells by a subject, the method comprising administering to the subject an effective amount of a pharmaceutical composition comprising [[a]] <u>an isolated</u> D isomer of an inhibitor of indoleamine-2,3-dioxygenase, <u>wherein the inhibitor of indoleamine-2,3-dioxygenase is selected from the group consisting of 1-methyl-D-tryptophan, β-(3-benzofuranyl)-D-alanine, β-(3-benzo(b)thienyl)-D-alanine, 6-nitro-D-tryptophan, and combinations thereof.</u>
- 2. (Currently amended) [[The]] A method of delaying the relapse or progression of a tumor in a subject: the method comprising administering an effective amount of a pharmaceutical composition comprising an isolated D isomer of an inhibitor of indoleamine-2,3-dioxygenase, claim 42 wherein the inhibitor of indoleamine-2,3-dioxygenase is selected from the group consisting of 1-methyl-D-tryptophan, β -(3-benzofuranyl)-D-alanine, β -(3-benzo(b)thienyl)-D-alanine, [[and]] 6-nitro-D-tryptophan and combinations thereof.
- 3. (Original) The method of claim 2 wherein the inhibitor of indoleamine-2,3-dioxygenase is 1-methyl-D-tryptophan.
- 4. (Cancel)

Page 3 of 14

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5. (Withdrawn/currently amended) The method of claim[[4]] 1, wherein the tumor cells are a cancer selected from the group consisting of melanoma, colon cancer, pancreatic cancer, breast cancer, prostate cancer, lung cancer, leukemia, brain tumors, lymphoma, sarcoma, ovarian cancer and Kaposi's sarcoma.

- 6. (Currently amended) The method of claim [[42]] 2 further comprising administering one or more chemotherapeutic agents to the subject.
- 7. (Original) The method of claim 6 wherein at least one chemotherapeutic agent is selected from the group consisting of cyclophosphamide, methotrexate, fluorouracil, doxorubicin, vincristine, ifosfamide, cisplatin, gemeytabine, busulfan, ara-C, and combinations thereof.
- 8. (Currently amended) The method of claim [[42]] 2 wherein the phannaceutical composition further comprises at least one chemotherapeutic agent.
- 9. (Original) The method of claim 8 wherein at least one chemotherapeutic agent is selected from the group consisting of cyclophosphamide, methotrexate, fluorouracil, doxorubicin, vincristine, ifosfamide, cisplatin, gemcytabine, busulfan, ara-C, and combinations thereof.
- 10. (Currently amended) The method of claim [[42]] 2 further comprising administering radiation therapy.

11-16. (Cancel)

17. (Currently amended) The method of claim [[42]] 2 wherein the pharmaceutical composition is administered in combination with a cytokine.

Page 4 of 14

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- 18. (Original) The method of claim 17 wherein the cytokine is granulocyte-macrophage colony stimulating factor (GM-CSF) or flt3-ligand.
- 19. (Currently amended) The method of claim [[42]] 2 wherein the pharmaceutical composition further comprises a cytokine.
- 20. (Currently amended) The method of claim [[42]] 2 wherein the pharmaceutical composition is administered in combination with a vaccine.
- 21. (Original) The method of claim 20, wherein the vaccine is a tumor vaccine.
- 22. (Original) The method of claim 21 wherein the tumor vaccine is a melanoma vaccine.
- 23. (Original) The method of claim 21 wherein the tumor vaccine comprises genetically modified tumor cells.
- 24. (Original) The method of claim 23 wherein the genetically modified tumor cells are transfected with granulocyte-macrophage stimulating factor (GM-CSF).
- 25. (Cancel)
- 26. (Original) The method of claim 21 wherein the tumor vaccine comprises dendritic cells.
- 27. (Withdrawn/currently amended) A method of stimulating an immune response to a tumor in a subject, the method comprising administering to the subject an effective amount of a pharmaceutical composition comprising [[a]] an isolated D isomer of an inhibitor of

Page 5 of 14

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indolearnine-2,3-dioxygenase, wherein the inhibitor of indolearnine-2,3-dioxygenase is selected from the group consisting of 1-methyl-D-tryptophan, β-(3-benzofuranyl)-D-alanine, β-(3-benzo(b)thienyl)-D-alanine, 6-nitro-D-tryptophan, and combinations thereof.

28-42. (Cancel)

43. (Withdrawn/currently amended) A method of treating a subject suffering from a neoplastic condition, the method comprising administering to the subject an effective amount of a pharmaceutical composition comprising [[a]] an isolated D isomer of an inhibitor of indoleamine-2,3-dioxygenase, wherein the inhibitor of indoleamine-2,3-dioxygenase is selected from the group consisting of 1-methyl-D-tryptophan, β-(3-benzofuranyl)-D-alanine, β-(3-benzofuran

44-47. (Cancel)